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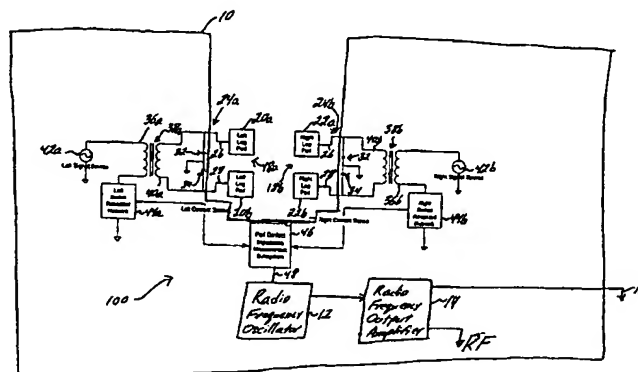
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(54) Title: MULTIPLE RF RETURN PAD CONTACT DETECTION SYSTEM



(57) Abstract: A multiple RF return pad contact detection system is provided which is adaptive to different physiological characteristics of patients without being susceptible to electro-surgical current interference (e.g., interference or measurement interaction between components of the detection system). The detection system can measure or sense the contact resistance or impedance between the patient and pairs of RF return pads or return electrodes where multiple pairs of RF return pads are utilized due to the high current frequently needed during electro-surgery while eliminating or minimizing the risk of measurement interaction between the RF return pad pairs. The system allows for the independent and simultaneous measurement of the pad contact impedance for each pair of RF return pads. If the impedance of any pad pair is above a predetermined limit, the system turns off or reduces the electro-surgical output of the electro-surgical generator to prevent excess heating. The system eliminates or minimizes interference or measurement interaction between the pad pairs by providing a different signal source frequency for each pad contact pair, but a frequency which matches an associated series resonant network frequency. The current that flows in the series resonant network is a direct reflection or function of the pad impedance of the corresponding pad pair.

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## **MULTIPLE RF RETURN PAD CONTACT DETECTION SYSTEM**

### **BACKGROUND**

#### **1. Technical Field**

The present disclosure is directed to electrosurgery and, in particular, to circuitry for measuring or sensing the contact resistance or impedance between the patient and pairs of RF return pad contacts or electrodes employed in such surgery.

#### **2. Description of the Related Art**

One potential risk involved in electrosurgery is the possibility of stray electrical currents causing excess heating proximate the RF return pad contacts or patient return electrodes. The most common conditions which are thought to lead to excess heating include:

(1) Tenting: Lifting of the return electrode from the patient due to patient movement or improper application. This situation may lead to excess heating if the area of electrode-patient contact is significantly reduced;

(2) Incorrect Application Site: Application of a return electrode over a highly resistive body location (e.g., excessive adipose tissue, scar tissue, erythema or lesions, excessive hair) will lead to a greater, more rapid temperature increase. Or, if the electrode is not applied to the patient (i.e. electrode hangs freely or is attached to another surface), the current may seek an alternate return path such as the table or monitoring electrodes; and

(3) Gel drying either due to premature opening of the electrode pouch or use of an electrode which has exceeded the recommended shelf life.

Many monitor or detection systems have been developed in the past, but most cannot directly guard against all three of the above listed situations. In order to protect against these potentially hazardous situations, the contact resistance or impedance between the return electrode and the patient should be monitored in addition to the continuity of the patient return circuit.

Safety circuitry is known whereby split (or double) patient electrodes are employed and a DC current (see German Pat. No. 1,139,927, published Nov. 22, 1962) or an AC current (see U.S. Pat. Nos. 3,933,157 and 4,200,104) is passed between the split electrodes to sense the contact resistance or impedance between the patient and the electrodes. U.S. Pat. No. 3,913,583 discloses circuitry for reducing the current passing through the patient depending upon the area of contact of the patient with a solid, patient plate. A saturable reactor is

included in the output circuit, the impedance of which varies depending upon the sensed impedance of the contact area.

The above systems are subject to at least one or more of the following shortcomings:

- (a) lack of sensitivity or adaptiveness to different physiological characteristics of patients and
- 5 (b) susceptibility to electrosurgical current interference when monitoring is continued during electrosurgical activation.

U.S. Pat. Nos. 4,416,276 and 4,416,277 describe a split-patient return electrode monitoring system which is adaptive to different physiological characteristics of patients, and a return electrode monitoring system which has little, if any, susceptibility to electrosurgical current interference when monitoring is continued during electrosurgical activation. The  
10 entire contents of both U.S. Pat. Nos. 4,416,276 and 4,416,277 are incorporated herein by reference.

Still a need exists for a detection or monitoring system, which is: 1) adaptive to different physiological characteristics of patients; 2) has little, if any, susceptibility to  
15 electrosurgical current interference, (including interference or measurement interaction between components of the detection system); 3) can measure or sense the contact resistance or impedance between the patient and pairs of RF return pads or electrodes where multiple pairs of RF return pads are utilized due to the high current frequently needed during electrosurgery, such as during tissue ablation; and 4) eliminates or minimizes the risk of  
20 measurement interaction between the RF return pad pairs.

Therefore, it is an aspect of the invention to provide a multiple RF return pad contact detection system for use during electrosurgical activation which achieves the above objectives.

### SUMMARY

25 A multiple RF return pad contact detection system is disclosed which is adaptive to different physiological characteristics of patients, without being susceptible to electrosurgical current interference. The detection system includes interference or measurement interaction between components of the detection system which can measure or sense the contact resistance or impedance between the patient and pairs of RF return pads or electrodes when  
30 multiple pairs of RF return pads are utilized. Due to the high current frequently needed during electrosurgery, such as during tissue ablation, the detection system advantageously eliminates or minimizes the risk of measurement interaction between the RF return pad pairs.

The circuitry of the multiple RF return pad contact detection system is preferably provided within an electrosurgical generator for controlling the generator according to various measurements, such as the contact resistance or impedance between the patient and pairs of RF return pads or return electrodes. Advantageously, the system allows for the independent and simultaneous measurement of the pad contact impedance for each pair of RF return pads. If the impedance of any pad pair is above a predetermined limit, the system advantageously turns off or reduces the electrosurgical output of the electrosurgical generator to prevent excess heating.

The system eliminates or minimizes interference or measurement interaction between the pad pairs by providing a different signal source frequency for each pad contact pair, but a frequency which matches an associated series resonant network frequency. The current that flows in the series resonant network is a direct reflection or function of the pad impedance of the corresponding pad pair. Since the two resonant networks are tuned to different frequencies, there is minimal interaction, if any, within the system, which advantageously reduces the chances of inaccurate measurements.

The system may advantageously include or be modified to include a multiplexer to multiplex the measurements corresponding to each pad contact pair to eliminate or minimize measurement interaction and also minimize hardware resources.

More specifically, the present disclosure relates to a return pad contact detection system for use with at least two pairs of patient return pads adapted for contacting a patient. Each pair of the at least two pairs of patient return pads has two conductors attached to a corresponding patient return pad for connecting the pad to a source of a.c. energy passing through the pad. The return pad contact detection system includes: at least two signal sources for generating an operating current for a corresponding pair of the at least two pairs of patient return pads; means for applying the operating current through the conductors to the at least two pairs of patient return pads; and at least two resonant circuits each corresponding to at least one pair of patient return pads and one of the at least two signal sources. Each of the at least two resonant circuits responsive to the current for producing a signal which is a function of the impedance between the two corresponding patient return pads. The at least two resonant circuits are tuned to different frequencies and are tuned to the substantially the same frequency as the corresponding signal source for substantially minimizing measurement interaction between the at least two pairs of patient return pads when the operating currents are simultaneously applied to the corresponding pairs of patient return pads. As can be

appreciated from the present disclosure, in one embodiment the frequency of the electrosurgical current may be substantially different from that of the operating current.

Preferably, the at least two resonant circuits are RCL series resonant circuits having minimum impedance at the resonant frequency. Advantageously, the system may include means for establishing a desired range having at least an upper limit for the impedance the system may also include determining means responsive to said signal for determining whether the impedance is within the desired range. Preferably, the means for establishing a desired range includes means for generating a reference signal corresponding to the upper limit and wherein the determining means includes comparator means for comparing the signal which is a function of the impedance with the reference signal.

Advantageously, the system may further include means for generating a control signal for controlling the operation of the electrosurgical generator according to the determination made by the comparator means.

The desired range may advantageously include a lower limit for the impedance and wherein the means for establishing a desired range includes means for generating a reference signal corresponding to the lower limit and wherein the determining means includes comparator means for comparing the signal which is a function of the impedance with the reference signal. Preferably, the lower limit for the impedance is about 20 ohms and the upper limit for the impedance is about 144 ohms.

Advantageously, the means for applying the operating current includes at least two transformers each for coupling the corresponding pair of patient return pads to the corresponding signal source. The secondary winding of each transformer is connected to the corresponding pair of patient return pads and the primary winding thereof is in circuit with the corresponding signal source and resonant circuit.

The present disclosure also relates to a return pad contact detection system for use with at least two pairs of patient return pads adapted for contacting a patient. Each pair of the at least two pairs of patient return pads having two conductors attached to a corresponding patient return pad for connecting the pad to a source of a.c. energy passing through the pad. The return pad contact detection system includes: means for producing and applying a corresponding current signal for each of the at least two pairs of patient return pads; resonant circuitry means responsive to the impedance between the at least two pairs of patient return pads for producing at least one signal which is a function of the impedance; and measurement means for receiving the at least one signal for determining the impedance between the at least

two pairs of patient return pads. Advantageously, the frequency of the corresponding current signal for each of the at least two pairs of patient return pads is substantially equal to the frequency of at least a portion of the resonant circuitry means.

Preferably, the means for producing and applying the corresponding current signal  
5 includes at least two a.c. signal sources. Advantageously, the resonant circuitry means includes at least two RCL series resonant circuits having minimum impedance at the resonant frequency.

The system may advantageously further include: means for establishing a desired range having at least an upper limit for the impedance and determining means responsive to  
10 the at least one signal for determining whether the impedance is within the desired range. Much like the embodiment above, the means for establishing a desired range includes means for generating a reference signal corresponding to the upper limit and wherein the determining means includes comparator means for comparing the at least one signal which is a function of the impedance with the reference signal. Preferably, the system further  
15 includes means for generating a control signal for controlling the operation of the electrosurgical generator according to the determination made by the comparator means.

Advantageously, the desired range includes a lower limit for the impedance and wherein the means for establishing a desired range includes means for generating a reference signal corresponding to the lower limit and wherein the determining means includes  
20 comparator means for comparing the at least one signal which is a function of the impedance with the reference signal. Preferably, the lower limit for said impedance is about 20 ohms and the upper limit for said impedance is about 144 ohms.

Advantageously, the means for producing and applying the corresponding current signal includes at least two transformers wherein the secondary winding of each transformer  
25 is connected to a corresponding pair of patient return pads and the primary winding thereof is in circuit with a portion of the means for applying and producing the corresponding current signal and the portion of the resonant circuitry means.

Advantageously the system may include two capacitors connected in series connected in parallel with the secondary winding of each transformer.

Further features of the multiple RF return pad contact detection system of the  
30 invention will become more readily apparent to those skilled in the art from the following detailed description of the apparatus taken in conjunction with the drawing.

### **BRIEF DESCRIPTION OF THE DRAWINGS**

Various embodiments of the invention will be described herein below with reference to the drawings wherein:

5        FIG. 1 is a schematic diagram of the multiple RF return pad contact detection system in accordance with a preferred embodiment of the invention; and

      FIG. 2 is a graph illustrating the operation of the pad contact impedance measurement subsystem of FIG. 1.

### **DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS**

10        Reference should be made to the drawings where like reference numerals refer to similar elements. Referring to FIG. 1, there is shown a schematic diagram of the multiple RF return pad contact detection system 100 of the present invention wherein electrosurgical generator 10 includes known circuitry such as a radio frequency oscillator 12 and an output amplifier 14 which generate an electrosurgical current. This current is applied to a patient  
15        (not shown) via an active electrode 16. The electrosurgical current is returned to the generator 10 via pad contact pairs or return electrode pairs 18a, 18b having pads or electrodes 20a, 20b and 22a, 22b and a corresponding two conductor patient cable 24a, 24b having leads 26 and 28. Two capacitors 32 and 34 are connected across each of the secondary windings 40a, 40b of transformer 38a, 38b.

20        Each primary winding 36a, 36b is connected to a corresponding a.c. signal source 42a, 42b and a series resonant network 44a, 44b. The purpose of each series resonant network 44a, 44b is to produce a current (i.e., left and right current senses) which is a function of the impedance between pads or electrodes 20a, 20b and 22a, 22b.

      The system 100 eliminates or minimizes interference or measurement interaction  
25        between the pads 20a, 20b and 22a, 22b, while allowing for the independent and simultaneous measurement of the pad contact impedance for each pair of RF return pads by having each a.c. signal source 42a, 42b provide a different signal source frequency for its corresponding pad contact pair. The frequency of each series resonant network 44a, 44b is tuned to match the frequency of the current produced by its associated a.c. signal source 42a,  
30        42b.

      Accordingly, the frequency of one of the series resonant networks 44a is different from the frequency of the other series resonant network 44b. Hence, there is minimal

interaction, if any, between the left and right circuitry of the system 100, especially the two contact pad pairs 18a, 18b. This essentially eliminates inaccurate or confusing measurements.

Additionally, the frequency of the electrosurgical current produced by the electrosurgical generator 10 is substantially different from that of the current produced by the a.c. signal sources 42a, 42b.

The current that flows in each series resonant network 44a, 44b, i.e., left and right current senses, is a direct reflection or function of the pad impedance of the corresponding pad contact pair 18a, 18b according to the physics of a series resonant network. Each series resonant network 44a, 44b is an RCL network or a combination of R (resistance), L (inductance) and C (capacitance). In a preferred embodiment of the series resonant networks 44a, 44b, the inductive component for each network is integrated into the respective transformer 38a, 38b.

The frequency response of a series resonant network has a maximum resonant frequency  $f_R$ . At the resonant frequency, the series resonant network has the minimum impedance, as opposed to a parallel resonant network which has the maximum impedance at the resonant frequency, and the phase angle is equal to zero degrees. The total impedance of a series resonant network is  $Z_T = jX_L - jX_C = R + j(X_L - X_C)$ . At resonance:  $X_L = X_C$ ,  $f_R = 1/(2\pi\sqrt{LC})$ ,  $Z_T = R$ , and  $V_L = V_C$ . The resonance of a series resonant network occurs when the inductive and capacitive reactances are equal in magnitude but cancel each other because they are 180 degrees apart in phase.

The left and right current senses are applied to pad contact impedance measurement subsystem 46 which determines whether the impedance measurements between pads or return electrodes 20a, 20b and 22a, 22b are within a desired range. The range is preferably adaptable to the physiological characteristics of the patient. If at least one of the impedance measurements is not within a desired range, an inhibit signal is applied over a line 48 to internally disable the electrosurgical generator 10 (or reduce the RF output therefrom) to prevent excess heating.

U.S. Pat. Nos. 4,416,276 and 4,416,277 describe a method for determining the desired range according to the physiological characteristics of the patient, the entire contents of these patents is incorporated herein by reference.

Preferably, the desired range for which the impedance must fall between return electrodes 20a, 20b and 22a, 22b is about 20 to about 144 ohms. If not, the electrosurgical generator 10 is disabled. Thus, in one method of operation of the present invention, the lower

limit is fixed at the nominal value of 20 ohms, thus reducing the onset of patient injury as a result of stray current paths which may surface if a contact pad or electrode is applied to a surface other than the patient. The upper limit is set to avoid such problems as those mentioned hereinbefore, i.e., tenting, incorrect application site, gel drying, etc.

5 In accordance with an important aspect of the invention, the upper limit is adjustable from the absolute maximum (typically about 144 ohms) downward to as low as typically 20 ohms to thereby provide for automatic adaptiveness to the physiological characteristics of the patient. This provides the multiple RF return pad contact detection system 100 of the present invention with significantly more control over the integrity of the RF pad contact or electrode  
10 connections without limiting the range of patient types with which the multiple RF return pad contact detection system 100 may be used or burdening the operator with additional concerns.

That is, the physiological characteristics can vary significantly from patient to patient and from one location site for the pad pairs to another. Thus, patients may vary in their respective amounts of adipose tissue (which is one determining factor in the impedance  
15 measurement between the various pads) without effecting the detection system. Further, for a particular patient, one location site may be more fatty, hairy or scarred than another. Again, this does not reduce the effectiveness of the system, i.e., all of these factors typically affect the impedance measured between pads 20a, 20b and 22a, 22b and thus concern the operator as to which site is optimal for a particular patient. Such concerns are eliminated in  
20 accordance with the present invention by providing for automatic adaptability to the physiological characteristics of the patient.

Reference should now be made to FIG. 2 which is a graph illustrating the operation of pad contact impedance measurement subsystem 46.

During operation, the desired impedance range (that is, the acceptable range of the  
25 impedance detected between pads 20a, 20b and 22a, 22b) is preset when the power is turned on to an upper limit of, for example, 120 ohms and a lower limit of, for example, 20 ohms as can be seen at time  $T=0$  seconds in FIG. 2. If the monitored impedance for any pad contact pair is determined to be outside of this range ( $T=A$  seconds) by comparing the current sense signal (or a signal derived there from) with a reference signal (e.g., a signal equal to 120 ohms  
30 or 20 ohms) using comparator circuitry (e.g., when a pad pair or any single contact pad is not affixed to the patient) an alert will be asserted and the electrosurgical generator 10 will be disabled over line 48.

The impedance between two contact pads of a contact pad pair at any instant is designated the return RF electrode monitor (REM) Instantaneous Value (RIV) in FIG. 2. When the REM impedance enters the range ( $T=B$  seconds) bounded by the Upper Limit (UL) and the Lower Limit (LL), a timing sequence begins. If after five seconds the RIV is still  
5 within range ( $T=C$  seconds), the alert condition will cease and the REM impedance value is stored in memory. This is designated as REM Nominal Value (RNV). The upper limit is then reestablished as 120% of this amount. The 80 ohm RIV shown in FIG. 2 causes the upper limit to be at 96 ohms. This feature of the invention is particularly important because it is at this time ( $T=C$  seconds) that adaptation is initially made to the physiological  
10 characteristics of the patient. Note if the RIV were to exceed 96 ohms at a time between  $T=C$  and  $T=F$  seconds (while the upper limit is 96 ohms), the alert will be asserted and the electrosurgical generator 10 disabled.

However, if the upper limit had not been adjusted to 96 ohms, the alert would not have been asserted until after the RIV exceeded the initial 120 ohms upper limit as  
15 determined by the comparator circuitry, thus possibly heating one or both of the pads 20a, 20b and 22a, 22b. This situation is of course exacerbated if the patient's initial RIV within the preset 20 to 120 ohm range is 30 ohms.

An initial RIV of 10 ohms within the preset range of 20 to 120 ohms sets an upper limit of 144 ohms.

20 In accordance with another aspect of the invention, it has been observed that the impedance between contact pads of contact pad pairs decreases over a relatively long period, such as a number of hours. Since many surgical procedures can extend a number of hours, this effect is also taken into consideration in the present invention. Accordingly, RIV is continuously monitored and any minima in REM impedance (e.g., a downward trend  
25 followed by a constant or upward trend in REM impedance) initiates a new five second timing interval ( $T=E$  seconds) at the end of which the RNV is updated to the RIV if the RIV is lower ( $T=F$  seconds). The REM upper limit of 120% of RNV is re-established at this time. The five second interval causes any temporary negative change in REM impedance ( $T=D$  seconds) to be disregarded. Operation will continue in this manner provided RIV does not  
30 exceed the upper limit of 120% RNV or drop below the lower limit of 20 ohms. Exceeding the upper limit ( $T=G$  seconds) causes an alert and the electrosurgical generator 10 is disabled. It will remain in alert until the RIV drops to 115% of RNV or less ( $T=H$  seconds) or until the system 100 is reinitialized. RIV dropping to less than 20 ohms ( $T=I$  seconds) causes a similar

alert which continues until either the RIV exceeds 24 ohms ( $T=J$  seconds) or the system 100 is reinitialized. The hysteresis in the limits of the REM range (that is, the changing of the upper limit to 115% of RNV and the lower limit to 24 ohms in the previous examples) prevents erratic alerting when RIV is marginal.

5 It should be noted in the example of FIG. 2 that the alert actually does not turn off when RIV returns to a value greater than 24 ohms because the pad pairs are removed before 5 seconds after  $T=J$  seconds elapse. Thus, the alarm stays on due to the removal of the pad contact pairs 18a, 18b.

10 Removing the pad contact pairs 18a, 18b from the patient or unplugging the cables 26, 28 from the electrosurgical generator 10 ( $T=K$  seconds) for more than one second causes the system 100 to be reinitialized to the original limits of 120 and 20 ohms. This permits a pad to be relocated or replaced ( $T=L$  seconds) without switching the electrosurgical generator 10 off. The RIV at the new location is 110 ohms and 120% RNV is 132 ohms. Thus, as described above, this is the one time (whenever RIV enters the 20 to 120 ohms range (either as preset  
15 during power on or as reinitialized as at  $T=K$  seconds) for the first time) that the upper limit can be raised during the normal REM cycle. Otherwise, it is continually decreased to adapt to the decreasing RIV impedance with the passage of time.

The preferred implementation of the foregoing FIG. 2 operation of the pad contact impedance measurement subsystem 46 is effected by a set of programmable instructions  
20 configured for execution by a microprocessor.

The system 100 could be modified by providing a multiplexer to multiplex the measurements corresponding to each pad contact pair 18a, 18b to eliminate or minimize measurement interaction and also minimize hardware resources.

25 Other pad contact pair arrangements can be provided in the system 100 of the present invention besides the pad pair arrangements shown in FIG. 1. For example, ten pad contact pairs 18 can be provided and connected to electrosurgical generator 10 by cables 26 and 28, where the corresponding a.c. signal source 42 and series resonant network 44 corresponding to each pad contact pair 18 are tuned to the same frequency which is different from the frequency of the other a.c. signal sources 42 and series resonant networks 44.

30 It is provided that the system 100 of the present invention allows for impedance comparisons to be performed between pad pairs. Therefore, if the pad pairs are placed symmetrically on the patient, i.e., left leg and right leg, comparison of the contact impedance can provide another degree of detection and safety.

Although the subject apparatus has been described with respect to preferred embodiments, it will be readily apparent to those having ordinary skill in the art to which it appertains that changes and modifications may be made thereto without departing from the spirit or scope of the subject apparatus.

**WHAT IS CLAIMED IS:**

1. A return pad contact detection system for use with at least two pairs of patient return pads adapted for contacting a patient, each pair of said at least two pairs of patient return pads having two conductors attached to a corresponding patient return pad for  
5 connecting the pad to a source of a.c. energy passing through the pad, said return pad contact detection system comprising:

at least two signal sources for generating an operating current for a corresponding pair of said at least two pairs of patient return pads;

means for applying the operating current through said conductors to said at least two  
10 pairs of patient return pads; and

at least two resonant circuits each corresponding to at least one pair of patient return pads and one of said at least two signal sources, each of said at least two resonant circuits responsive to said current for producing a signal which is a function of the impedance  
15 between said two corresponding patient return pads, said at least two resonant circuits being tuned to different frequencies and being tuned to the substantially the same frequency as the corresponding signal source for substantially minimizing measurement interaction between the at least two pairs of patient return pads when said operating currents are simultaneously applied to said corresponding pairs of patient return pads.

20 2. A system as in Claim 1, wherein said at least two resonant circuits are RCL series resonant circuits having minimum impedance at the resonant frequency.

3. A system as in any preceding claim, further comprising:

means for establishing a desired range having at least an upper limit for said  
25 impedance; and

determining means responsive to said signal for determining whether said impedance is within said desired range.

4. A system as in any preceding claim, wherein said means for establishing a  
30 desired range includes means for generating a reference signal corresponding to the upper limit and wherein said determining means includes comparator means for comparing the signal which is a function of said impedance with the reference signal.

5. A system as in any preceding claim, wherein said system further comprises means for generating a control signal for controlling the operation of said electrosurgical generator according to the determination made by said comparator means.

5 6. A system as in any preceding claim, wherein said desired range includes a lower limit for said impedance and wherein said means for establishing a desired range includes means for generating a reference signal corresponding to the lower limit and wherein said determining means includes comparator means for comparing the signal which is a function of said impedance with the reference signal.

10 7. A system as in any preceding claim, wherein said system further comprises means for generating a control signal for controlling the operation of said electrosurgical generator according to the determination made by said comparator means.

15 8. A system as in any preceding claim, wherein the lower limit for said impedance is about 20 ohms and the upper limit for said impedance is about 144 ohms.

9. A system as in any preceding claim, wherein said means for applying the operating current includes at least two transformers each for coupling the corresponding pair of patient return pads to the corresponding signal source, the secondary winding of each transformer being connected to the corresponding pair of patient return pads and the primary winding thereof being in circuit with the corresponding signal source and resonant circuit.

20 10. A system as in any preceding claim, wherein the frequency of said electrosurgical current is substantially different from that of said operating current.

11. A return pad contact detection system for use with at least two pairs of patient return pads adapted for contacting a patient, each pair of said at least two pairs of patient return pads having two conductors attached to a corresponding patient return pad for connecting the pad to a source of a.c. energy passing through the pad, said return pad contact detection system comprising:

30 means for producing and applying a corresponding current signal for each of said at least two pairs of patient return pads;

resonant circuitry means responsive to the impedance between said at least two pairs of patient return pads for producing at least one signal which is a function of said impedance; and

5 measurement means for receiving said at least one signal for determining the impedance between said at least two pairs of patient return pads, wherein the frequency of said corresponding current signal for each of said at least two pairs of patient return pads is substantially equal to the frequency of at least a portion of said resonant circuitry means.

12. A system as in Claim 11, wherein the means for producing and applying said  
10 corresponding current signal includes at least two a.c. signal sources.

13. A system as in Claim 11 or 12 wherein said resonant circuitry means includes at least two RCL series resonant circuits having minimum impedance at the resonant frequency.  
15

14. A system as in Claim 11, 12 or 13 further comprising:  
means for establishing a desired range having at least an upper limit for said impedance; and  
determining means responsive to said at least one signal for determining whether said  
20 impedance is within said desired range.

15. A system as in Claim 11, 12, 13 or 14 wherein said means for establishing a desired range includes means for generating a reference signal corresponding to the upper limit and wherein said determining means includes comparator means for comparing said at  
25 least one signal which is a function of said impedance with the reference signal.

16. A system as in Claim 11, 12, 13, 14 or 15 wherein said system further comprises means for generating a control signal for controlling the operation of said electrosurgical generator according to the determination made by said comparator means.  
30

17. A system as in Claim 11, 12, 13 or 14, wherein said desired range includes a lower limit for said impedance and wherein said means for establishing a desired range includes means for generating a reference signal corresponding to the lower limit and wherein

said determining means includes comparator means for comparing said at least one signal which is a function of said impedance with the reference signal.

18. A system as in Claim 11, 12, 13, 14, 15, 16 or 17, wherein said system further  
5 comprises means for generating a control signal for controlling the operation of said  
electrosurgical generator according to the determination made by said comparator means.

19. A system as in Claim 11, 12, 13, 14, 15, 16, 17 or 18 wherein the lower limit  
for said impedance is about 20 ohms and the upper limit for said impedance is about 144  
10 ohms.

20. A system as in Claim 11, 12, 13, 14, 15, 16, 17, 18 or 19 wherein said means  
for producing and applying said corresponding current signal includes at least two  
transformers, the secondary winding of each transformer being connected to a corresponding  
15 pair of patient return pads and the primary winding thereof being in circuit with a portion of  
the means for applying and producing the corresponding current signal and the portion of said  
resonant circuitry means.

21. A system as in Claim 11, 12, 13, 14, 15, 16, 17, 18, 19 or 20 further  
20 comprising two capacitors connected in series connected in parallel with the secondary  
winding of each transformer.

22. A system as in Claim 11, 12, 13, 14, 15, 16, 17, 18, 19, 20 or 21 wherein the  
frequency of said electrosurgical current is substantially different from that of said  
25 corresponding current signal.

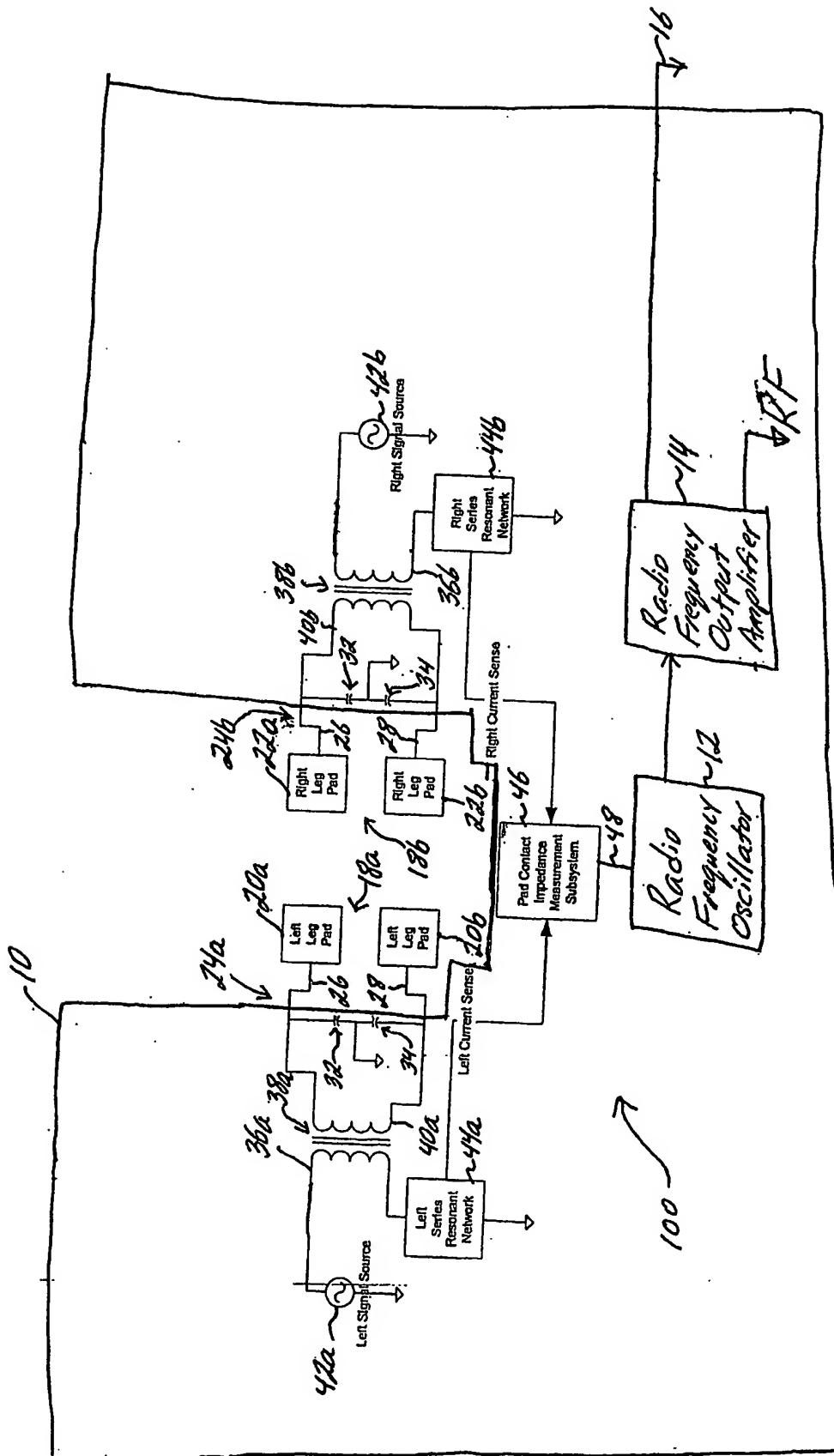
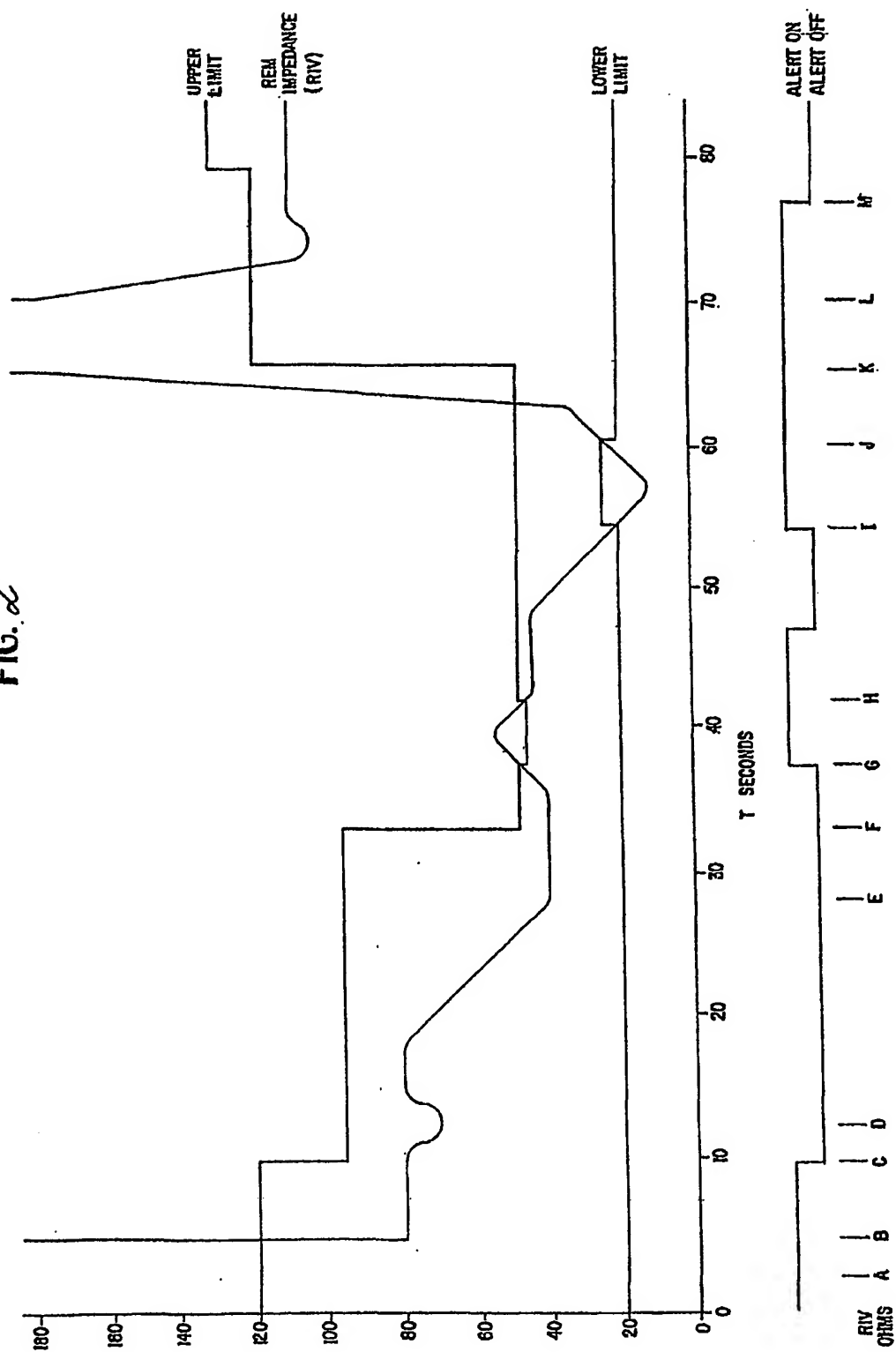


Fig. 1

FIG. 2



# INTERNATIONAL SEARCH REPORT

Intern: Application No  
PCT/US 03/28537

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 7 A61B18/12

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4 416 276 A (NEWTON DAVID W ET AL) 22 November 1983 (1983-11-22) cited in the application column 3, line 6 - column 4, line 15; figure 3	1,11
A	US 5 196 008 A (HAGEN UWE ET AL) 23 March 1993 (1993-03-23) abstract; figure 1	1,11
A	US 5 830 212 A (CARTMELL ROBERT LOUIS ET AL) 3 November 1998 (1998-11-03) column 7, line 16 - line 34; figure 4	1,11
A	US 6 275 786 B1 (DANERS FELIX) 14 August 2001 (2001-08-14) column 3, line 44 - line 67; figure 1	1,11
	-/--	

☒ Further documents are listed in the continuation of box C.

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Date of the actual completion of the international search

12 December 2003

Date of mailing of the international search report

19/12/2003

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>EP 0 836 868 A (BERCHTOLD GMBH &amp; CO GEB)  22 April 1998 (1998-04-22)  column 4, line 36 -column 5, line 14;  figure 1</p> <p>-----</p>	1,11

# INTERNATIONAL SEARCH REPORT

Internal Application No  
PCT/US 03/28537

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 4416276	A	22-11-1983	AU 541482 B2	10-01-1985
			AU 8941482 A	05-05-1983
			CA 1200286 A1	04-02-1986
			DE 3239640 A1	19-05-1983
			DE 3249766 C2	18-08-1988
			JP 1076846 A	22-03-1989
			JP 7073585 B	09-08-1995
			JP 1871580 C	06-09-1994
			JP 3049577 B	30-07-1991
			JP 58103445 A	20-06-1983
US 5196008	A	23-03-1993	EP 0416158 A1	13-03-1991
			AT 114441 T	15-12-1994
			AU 635964 B2	08-04-1993
			AU 6224390 A	14-03-1991
			CA 2024660 A1	08-03-1991
			DE 58908704 D1	12-01-1995
			ES 2064404 T3	01-02-1995
			JP 3106358 A	02-05-1991
US 5830212	A	03-11-1998	NONE	
US 6275786	B1	14-08-2001	DE 19714972 A1	22-10-1998
			WO 9844855 A1	15-10-1998
			EP 1115343 A1	18-07-2001
EP 0836868	A	22-04-1998	DE 19643127 A1	23-04-1998
			US 5971981 A	26-10-1999
			EP 0836868 A2	22-04-1998

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